





**GALVANIZED ALUMINIUM SHEET****Patent number:** WO9743467**Publication date:** 1997-11-20**Inventor:** GEHLHAAR HORST (DE); SPANJERS MARTINUS  
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MEER WILHELMUS JACOBUS V D (NL)**Classification:****- international:** C25D5/44; C25F3/04**- european:** C25D5/44, C25F3/04**Application number:** WO1997EP02329 19970507**Priority number(s):** NL19961003090 19960513**Also published as:** EP0904427 (A1)  
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In a method for applying a zinc layer onto an aluminium or aluminium alloy sheet, comprising pretreating the surface and applying the layer by electrolytic galvanizing, the pretreating comprises electrochemical graining of said surface, for example in a solution having a pH less than 3 with an alternating current applied between the sheet and an electrode. In an alternative method, the pretreating includes applying a preliminary zinc layer by immersing the surface in a zinc-containing alkaline solution, applying a potential to the sheet and reversing the polarity of said potential at least once. The sheets are useful as for example building cladding sheets and automotive panels.

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<b>(21) International Application Number:</b> PCT/EP97/02329 <b>(22) International Filing Date:</b> 7 May 1997 (07.05.97)  <b>(30) Priority Data:</b> 1003090 13 May 1996 (13.05.96) NL  <b>(71) Applicant (for all designated States except US):</b> HOOGOVENS ALUMINIUM BAUSYSTEME GMBH [DE/DE]; August-Horch-Strasse 20-22, D-56070 Koblenz (DE).  <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> GEHLHAAR, Horst [DE/DE]; Holbeinstrasse 2, D-56626 Andemach (DE); SPANJERS, Martinus, Godefridus, Johannes [NL/NL]; Zeewijkplein 108, NL-1974 PE IJmuiden (NL). MOOIJ, Joop, Nicolaas [NL/NL]; Walstro 56, NL-1902 JP Casticum (NL). VAN DER MEER, Wilhelmus, Jacobus [NL/NL]; Baroniestraat 15, NL-1079 PB Amsterdam (NL).  <b>(74) Agent:</b> HANSEN, Willem, Joseph, Maria; Hoogovens Corporate Services B.V., P.O. Box 10000, NL-1970 CA IJmuiden (NL).		<b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> GALVANIZED ALUMINIUM SHEET  <b>(57) Abstract</b>  In a method for applying a zinc layer onto an aluminium or aluminium alloy sheet, comprising pretreating the surface and applying the layer by electrolytic galvanizing, the pretreating comprises electrochemical graining of said surface, for example in a solution having a pH less than 3 with an alternating current applied between the sheet and an electrode. In an alternative method, the pretreating includes applying a preliminary zinc layer by immersing the surface in a zinc-containing alkaline solution, applying a potential to the sheet and reversing the polarity of said potential at least once. The sheets are useful as for example building cladding sheets and automotive panels.		

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## GALVANIZED ALUMINIUM SHEET

## TECHNICAL FIELD

The invention relates to methods for applying a layer comprising zinc onto at least one surface of an aluminium or aluminium alloy sheet, comprising in succession a pretreatment step and an electrolytic galvanizing step. The invention also relates to the galvanized aluminium or aluminium alloy sheet produced by the methods, and the use of such sheet in building structures and as automotive body sheet. In the following description, for brevity we refer to aluminium sheets, but this term is to be understood to include aluminium alloy sheets.

## BACKGROUND ART

Aluminium sheet is used on a wide scale in building structures as interior and/or exterior panels of buildings both for façade and roofing structures. An advantage of this is that the low specific weight of the aluminium means that the building structure may be made considerably lighter than for example with steel sheet. A disadvantage of untreated aluminium sheet is that the building structure reflects a large amount of light which limits its application in the immediate vicinity of airfields, for example. A solution to this problem is to provide the aluminium sheet with a surface layer, for example zinc, while retaining the structural advantages. An advantage of a galvanized aluminium sheet is that it reflects less light and has good corrosion resistance. A

further advantage of galvanized aluminium sheet is that the building structure made with it requires little maintenance due to the durability of the AlZn system. A further advantage of galvanized aluminium sheet is that the appearance of the aluminium sheet changes slowly over time, i.e. it "lives". This last property, the so-called patina effect, is much sought after by architects for application in buildings of their design. Galvanizing the aluminium sheet considerably increases the applicability of the aluminium sheet.

Methods of electrolytically plating zinc onto aluminium (galvanizing) are well known. To improve the applied zinc layer, pretreatments of the aluminium surface have been proposed.

JP-A-52005630 discloses electroplating one of Cu, Ni, Zn, Sn, Pb, Cd and Cr onto the chemically roughened surface of an aluminium or aluminium workpiece. The roughening is performed in two stages: first in alkali  $\text{pH} \geq 11$  or a fluoric acid solution, and second using a mixed mineral acid solution.

In EP-A-0497302 a pretreatment step consisting of degreasing and pickling is performed and cathodic zinc electroplating is then carried out in two steps, thereby forming two adjacent zinc layers. The layers may contain Ni or Fe additions to improve bonding.

Another method for pretreating and galvanizing an aluminium sheet is given in EP-A-0498436, which describes a method for the continuous electrolytic application of a

zinc layer onto an aluminium sheet intended for processing into an automotive body sheet, after which the aluminium sheet is provided with a paint layer. The method comprises in succession the steps (i) alkaline degreasing, (ii) pickling in an acid solution, (iii) anodizing and cathodic zinc electrolysis of the aluminium sheet in the same acid solution. In between the steps the aluminium sheet is cleaned by rinsing with water. The anodization and electrolysis process is not dependent on the extent of pretreatment, which may even be omitted. A disadvantage of galvanized aluminium sheet obtained by this method is that the bonding of the applied zinc layer on the aluminium sheet is very poor when the galvanized aluminium sheet is highly deformed, for example by bending.

Zinc plating of wrought aluminium sheet as a precursor to zinc phosphate treatment and painting is described in US-A-5176963. The zinc plating may comprise two stages, namely displacement plating from an alkaline bath and electroplating from an acid bath. There is no disclosure of the application of a potential to the sheet during the displacement plating stage.

#### DISCLOSURE OF THE INVENTION

An object of the invention is to provide a method for pretreating and galvanizing an aluminium sheet by which a very effective bond is obtained between the applied zinc layer and the aluminium sheet, the bond remaining effective under large subsequent deformation of

the sheet. An additional object of the invention is that the method for pretreating and galvanizing the aluminium sheet may be carried out as a continuous process.

According to the invention in one aspect there is provided a method for applying a layer comprising zinc onto at least one surface of an aluminium or aluminium alloy sheet, comprising the steps of pretreating said surface and applying the layer by electrolytic galvanizing, characterized in that the pretreating step comprises electrochemical graining of said surface.

Electrolytic graining is a process known in the art and is one example of the technique of graining. It is known in the art (see "Study of the mechanism of the A.C. electrolytic graining of aluminium" by P. Laevers, Brussels Free University, November 1995 and EP-A-586504) to use electrolytic graining for roughening of aluminium sheet, typically an aluminium-litho sheet, but the use of graining as a pretreatment in electrolytic galvanizing of aluminium sheet is novel.

By the electrochemical graining pretreatment step of the invention an effective bond between the electrolytically applied zinc layer and the aluminium sheet is formed, the bond remaining effective during subsequent deformation of the aluminium sheet, for example by bending. It can also achieve the effect that the resulting aluminium sheet has a very good corrosion resistance. Furthermore the method may be carried out in a continuous process.

The invention is based in part on the insight that to obtain a well-bonded zinc layer on the aluminium sheet so that the bond remains effective under great deformation of the galvanized aluminium sheet, the pretreatment is extremely important. It is believed that the electrochemical graining of the aluminium sheet produces a roughened surface so that the subsequently electrolytically applied zinc layer is also mechanically bonded. This mechanical bonding is partly responsible for achieving the effect that the zinc layer remains bonded under large deformation of the galvanized aluminium sheet.

Very good results, including effective bonding of the electrolytically applied zinc layer may be obtained with graining process parameters comprising one or more of:

- (a) alternating current, preferably with a frequency in the range from 10-1000 Hz, more preferably 40-100 Hz;
- (b) current density of 5-200 A/dm<sup>2</sup>, preferably 10-100 A/dm<sup>2</sup>, and more preferably 20-50 A/dm<sup>2</sup>;
- (c) bath temperature 15-60°C, preferably 40-50°C;
- (d) treatment time 0.2-60 s, preferably 0.5-10 s, and more preferably 2-3 s;
- (e) bath acidity pH < 3, and preferably pH < 2.

A further advantage is that these process parameters lend themselves to application in a continuous process operation.



As mentioned, the electrolytic graining is preferably performed in an acid solution. Various acids may be used for this, but preferably use is made of a hydrochloric acid solution or a nitric acid solution. The graining step may be a part of a pretreatment cycle. A typical preferred pretreatment cycle comprises in succession:

- (1) alkaline degreasing;
- (2) electrolyte graining;
- (3) anodizing e.g. in a sulphate solution;

and may also comprise rinsing with water between these steps. However, the method is not limited to this pretreatment cycle. Better alkaline degreasing is achieved if simultaneously a direct current is applied in a range 1-20 A/dm<sup>2</sup>, preferably 5-15 A/dm<sup>2</sup>.

Preferably anodizing is carried out after the electrolytic graining.

In a further aspect, the invention consists in an aluminium or aluminium alloy sheet having an electrolytically applied layer comprising zinc in which said layer is adjacent to an aluminium or aluminium alloy surface roughened by electrochemical graining. By microscopic inspection of the aluminium-zinc interface it will be possible to detect the electrolytic graining.

In a second method, the invention provides a method for applying a layer comprising zinc onto at least one surface of an aluminium or aluminium alloy sheet, comprising the steps of pretreating said surface and

applying said layer by electrolytic galvanizing, said  
pretreating step including applying a preliminary layer  
comprising zinc onto said surface, characterized in that  
the application of said preliminary layer comprises (i)  
5 immersing said surface in a zinc-containing alkaline  
solution, (ii) applying a potential to the sheet to cause  
an electrolytic current to flow, and (iii) reversing the  
polarity of said potential at least once. This method  
may be combined with the electrolytic graining  
10 pretreatment described above.

By this method of applying a preliminary zinc  
layer, a very effective bond of the subsequent  
electrolytically applied zinc layer may thereby be  
obtained. An advantage of this pretreatment is that the  
15 whole pretreatment may be carried out in an alkaline  
solution. Thus only a single waste flow, which is  
alkaline, may be formed which represents a logistical  
advantage. In addition the aluminium sheet can be  
directly degreased. The entire pretreatment cycle can  
20 therefore consist essentially of one pretreatment step.

Zincate treatments are known for applying  
conversion layers onto aluminium, for example as known  
from "Oppervlaktebehandelingen van aluminium" by T. van  
der Klis and J.W. du Mortier published by the Vereniging  
25 voor Oppervlaktetechnieken voor Materialen, Bilthoven,  
NL, 3rd edition 1992, pp 406-409. A basic composition  
for a zincate pickle comprises 40-50 g/l ZnO and 400-500  
g/l NaOH. However, a zincate treatment according to the

present method, besides applying a thin zinc layer onto the aluminium sheet, also activates the surface of the aluminium sheet so that the zinc layer applied electrolytically subsequently bonds better. The thickness of the layer comprising zinc produced in the pretreatment step is not of major importance, but may be in the range 0.1 to 0.5 g/m<sup>2</sup>, and the ultimate layer thickness of the zinc is essentially determined during the electrolytic galvanizing stage.

In this second method of the invention, the aluminium sheet may be anodized, e.g. in a sulphate solution, prior to the electrolytic galvanizing. This can achieve the effect that the electrolytically applied zinc layer bonds very effectively to the aluminium sheet.

In the second method of the invention, during the zincate treatment, the aluminium sheet changes polarity at least once, preferably at least twice. Thus the aluminium sheet may be initially anode-connected (A) for a given time, and then cathode-connected (C). This connection sequence may be designated A-C. The aluminium sheet is preferably anode-connected first so that a part of the oxide layer goes into solution, after which a thin zinc layer deposits onto the aluminium sheet at the time when the aluminium sheet is cathode-connected. However C-A connection is also possible. The final connection is preferably as a cathode. Preferably the aluminium sheet is connected at least A-C-A-C. The duration of maintenance of each polarity is here called the

electrolysis time per polarity, and is preferably at least 0.5 s.

Very good results may be obtained with zincate process parameters comprising one or more of:

- 5 (a) bath temperature 10-60°C, preferably 20-30°C;
- (b) current density of 0.5-20 A/dm<sup>2</sup>, preferably 2-10 A/dm<sup>2</sup>;
- (c) electrolysis time per polarity 0.5-10 s, preferably 1-4 s;
- 10 (d) total immersion time 1-30 s, preferably 2-10 s;
- (e) bath composition comprising 10-300 g/l NaOH and 2-40 g/l ZnO, preferably 50-150 g/l NaOH and 5-20 g/l ZnO.

15 An advantage of these process parameters is that the pretreatment process lends itself well to application in a continuous process operation.

Following the pretreatment according to both methods of the invention the aluminium sheet is electrolytically galvanized in an acid solution.

20 Conventional galvanizing techniques may be used, and many different processes are suitable. Good results may be obtained when the galvanizing process parameters comprise one or more of:

- 25 (a) direct current with a current density of 10-100 A/dm<sup>2</sup>, preferably 30-70 A/dm<sup>2</sup>;
- (b) bath temperature 20-70°C, preferably 40-60°C;
- (c) treatment time 5-90 s, preferably 10-40 s;

- (d) electrolyte composition comprising zinc sulphate with 30-200 g/l zinc, preferably 60-120 g/l;
- (e) acidity of the electrolyte  $\text{pH} < 4$ , preferably approximately  $\text{pH} 2.5$ .

5 Using these parameters, a well-bonded layer comprising essentially zinc may be applied to the pretreated aluminium sheet, the bonding remaining effective under a large deformation of the galvanized aluminium sheet. A further advantage is that it is possible to perform a continuous process. With a  $\text{pH}$  of approximately 2.5 it is possible to buffer the electrolyte, thereby making the electrolytic galvanizing process considerably more stable.

10 The electrolyte composition used is not limited to a composition comprising a sulphate solution, and for example a chloride solution may also be applied.

15 According to a preferred version of the first method of the invention, wherein the pretreatment comprises electrolytic graining followed by a brief anodization after which the aluminium sheet is electrolytically galvanized, desirably the same electrolyte liquid bath is not used for the anodizing as for the galvanizing. Working with a continuous process operation, but one in which the electrolytes for the anodizing and the galvanizing are separate, has the advantage that the two process conditions may be controlled independently. Thus for the anodizing process preferably a  $\text{pH} < 2$  is used, and for the electrolytic

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galvanizing preferably a pH of approximately 2.5.

Moreover, the electrolyte for the anodizing may comprise either zinc sulphate or an iron sulphate, and also such metals as Ni or Cu, while the electrolyte for the galvanizing is preferably zinc sulphate.

Both methods in accordance with the invention are preferably carried out in a continuous process, although it is possible to carry them out batchwise.

As mentioned, the methods in accordance with the invention are suitable for galvanizing aluminium sheets made of aluminium and a wide range of aluminium alloys, such as aluminium alloys of the 1xxx type, the 3xxx type, and the 6xxx type, but also of the 2xxx type and the 5xxx type (AA designations).

The zinc layer applied in the methods of the invention may be essentially a pure zinc layer or may be primarily zinc but including minor amounts of impurity elements or deliberately added elements, as is known in the art. Typically such impurity elements or added elements are present at less than 10%, more usually less than 5% by weight in the zinc layer.

Another advantage of the methods in accordance with the invention is that the galvanized aluminium sheet which is produced has an atmospheric corrosion resistance almost comparable to that of zinc sheet, which has been used for application in building structures. This enables structures to be made lighter while retaining the good corrosion resistance. Moreover, the total zinc

consumption when using galvanized aluminium sheet in building structures is considerably less than when using zinc sheets.

The invention also consists in the galvanized aluminium sheet produced by the methods according to the invention.

In a further aspect galvanized aluminium sheet, obtained by either method, may be used in building systems such as facade and roofing structures, and windowsill structures. Galvanized aluminium sheet suitable for application in building systems such as those known under the trade names KAL-ZIP and KAL-BAU may be obtained by the invention. In the case of application of the galvanized aluminium sheet in building structures, the aluminium sheet preferably comprises a weight per unit area of applied zinc in the range of 10-300 g/m<sup>2</sup>, and more preferably 30-100 g/m<sup>2</sup>.

In another aspect galvanized aluminium sheet, obtained by either method, may be used in shaping applications such as for the manufacture of automotive body parts by pressing. In the case of use of the galvanized aluminium sheet as automotive body sheet, the aluminium sheet preferably comprises a weight per unit area of applied zinc in the range 5-100 g/m<sup>2</sup>, and more preferably in the range 5-40 g/m<sup>2</sup>.

#### BEST MODES OF CARRYING OUT THE INVENTION

The invention will now be illustrated by several non-limitative examples.

Example 1

Aluminium sheets manufactured from an AA3004 alloy suitable for application in building structures were pretreated and electrolytically galvanized in different ways in batch processes. The bonding of the applied zinc layer was then tested by the so-called tape method. This is a rapid method for testing bonding. A piece of tape or self-adhesive tape, for example such as that used in offices, is stuck onto the galvanized aluminium sheet and then pulled off again by hand. A value assessment is then given to the bondability by allocating a number where: (1) = excellent, (2) = good, (3) = fair, (4) = poor, (5) = very poor. For value assessments (1) and (2) the bonding was also tested using the known zero bending test and the known lock-form test. For value assessments (3) to (5) these supplementary bond tests were omitted. The galvanized aluminium sheets with value assessments of (1) or (2) were also tested for durability in a corrosive, maritime industrial environment.

Table 1 gives the main process parameters used and the value assessment for the bond. Between the different steps of the pretreatment and the galvanizing the aluminium sheets were rinsed with distilled water for at least 5 s.

The degreasing in tests (1) to (14) was carried out using Percy 6340-29 (trade name) produced by Henkel Metall Chemicals, concentration 10 g/l, treatment time 3



s, bath temperature approximately 65°C, direct current with a current density of approximately 10 A/dm<sup>2</sup>.

The pickling in tests (1), (2), (9), (11) and (12) was carried out in an HCl environment (8% HCl solution), pH = 1, treatment time approximately 5 s. In tests (6) and (7) an 8% H<sub>2</sub>SO<sub>4</sub> solution was used and a treatment time of approximately 5 s.

The graining in tests (3) to (5) was carried out in 1% HCl solution having a pH of about 1, bath temperature approximately 40°C, alternating current of approximately 50 Hz, current density of approximately 50 A/dm<sup>2</sup> and variable treatment time.

The anodizing in tests (5), (7) to (9), (12) and (16) was carried out with direct current, current density approximately 50 A/dm<sup>2</sup>, polarity positive, treatment time approximately 5 s, a ZnSO<sub>4</sub>·H<sub>2</sub>O electrolyte comprising approximately 90 g/l zinc, pH = 1.8, bath temperature approximately 50°C. In tests (10) and (11) the electrolyte comprised sulphuric acid, other parameters were identical to test (5), except that the temperature was approximately 70°C. In tests (13) and (14) sulphuric acid and phosphoric acid respectively were used for the electrolyte, pH = 1.8, bath temperature approximately 60°C, stainless steel electrode material, alternating current, electrolysis time approximately 2 s, current density approximately 2 A/dm<sup>2</sup>, total immersion time approximately 10 s.

In tests (15) to (17) the pretreatment was carried out by connecting the aluminium sheet A-C-A-C, electrolysis time per polarity approximately 3 s, bath temperature approximately 20°C, current density approximately 5 A/dm<sup>2</sup>, immersion time approximately 10 s, composition of electrolyte 100 g/l NaOH and 10 g/l ZnO. The thickness of the zinc layers formed by this zincate treatment were in the range 0.1 to 0.5 g/m<sup>2</sup>. In test (16) the aluminium sheet was anodized in addition to that treatment.

In tests (1) to (16) the aluminium sheets were electrolytically galvanized in a zinc sulphate electrolyte with 90 g/l zinc, pH approximately 2, direct current with a current density of approximately 50 A/dm<sup>2</sup>, bath temperature approximately 50°C, immersion time approximately 20 s. In test (17) the pH was approximately 2.5, other parameters being identical to tests (1) to (16). The electrolytically applied zinc layer was approximately 35 g/m<sup>2</sup>.

It may be inferred from the results given in Table 1 that a pretreatment consisting of degreasing and pickling (tests 1, 2 and 6) is insufficient to obtain a well-bonded layer comprising zinc.

A pretreatment consisting of degreasing and anodizing (tests 8, 10, 13 and 14) is also insufficient to obtain a well-bonded layer comprising zinc.

The combination of in succession degreasing, pickling and anodizing (tests 7, 9, 11 and 12) was also

insufficient to obtain a well-bonded layer comprising zinc.

Very good results were obtained when the aluminium sheet was electrolytically grained following the degreasing (tests 3 to 5). The best results were achieved with the method in accordance with test (5), in which anodizing followed the graining.

Very good results were also obtained when the aluminium sheet was pretreated in an alkaline environment comprising zinc while being A-C-A-C connected (tests 15 to 17). Good results were also obtained if, following on from that, an anodization step carried out (test 16).

The galvanized aluminium sheets obtained with the method in accordance with tests (3), (5) and (15) to (17) were also subjected to the zero bending test and the lock-form test. In all cases the bond of the zinc layer and the aluminium sheet remained intact.

Galvanized aluminium sheets manufactured with the method in accordance with tests (3), (5) and (15) to (17) were locally greatly deformed in such a way that the galvanized aluminium sheets were comparable in shape to KAL-ZIP and KAL-BAU. These galvanized aluminium sheets were then tested for durability by means of the Atmospheric Building Corrosion Test as described by B. Boelen in the article "New Product Test: The Atmospheric Building Corrosion Test (ABC Test)", published on the occasion of the ECCA Autumn Congress in Brussels on 27-28 November 1995, and compared with

untreated sheets of AA3004 alloy and pure zinc. After 6 weeks in a maritime industrial surrounding there was no visible difference between galvanized aluminium and pure zinc (Reinzink), while the AA3004 alloy was badly corroded particularly by the chloride. Among other things it can be inferred from the results that galvanized aluminium sheet manufactured in accordance with the invention has a durability comparable to a sheet of pure zinc.

Table 1

Test	Degreasing		Pickling		Graining		Anodizing		Galvanizing	Bonding rating
	Temp ('C)	Time (s)	Solution	Temp ('C)	Solution	Time (s)	Solution	Time (s)	pH	
1	65	3	HCl	50	-	-	-	-	2	5
2	65	3	HCl	80	-	-	-	-	2	5
3	65	3	-	-	HCl	30	-	-	2	2
4	65	3	-	-	HCl	5	-	-	2	3
5	65	3	-	-	HCl	3	ZnSO <sub>4</sub>	5	2	1
6	65	3	H <sub>2</sub> SO <sub>4</sub>	70	-	-	-	-	2	5
7	65	3	H <sub>2</sub> SO <sub>4</sub>	70	-	-	ZnSO <sub>4</sub>	5	2	4
8	65	3	-	-	-	-	ZnSO <sub>4</sub>	5	2	4
9	65	3	HCl	50	-	-	ZnSO <sub>4</sub>	5	2	5
10	65	3	-	-	-	-	H <sub>2</sub> SO <sub>4</sub>	5	2	5
11	65	3	HCl	50	-	-	H <sub>2</sub> SO <sub>4</sub>	5	2	5
12	65	3	HCl	80	-	-	ZnSO <sub>4</sub>	5	2	5
13	65	3	-	-	-	-	H <sub>2</sub> SO <sub>4</sub>	2	2	4
14	65	3	-	-	-	-	H <sub>3</sub> PO <sub>4</sub>	2	2	4
15	Alkaline solution containing zinc + A-C-A-C connection						-	-	2	1
16							ZnSO <sub>4</sub>	5	2	1
17							-	-	2.5	1

Example 2

The process as described in test 5 of Example 1 was further performed in eight test runs on a continuous pilot line using AA3004 sheet material 0.26 m wide.

The process parameters used for the eight different test runs are listed in Table 2. In between

graining and anodizing also the sheet was rinsed with distilled water.

The galvanized sheet which was obtained was tested for the amount of zinc layer, and the bonding was tested using the zero bending test.

Degreasing was performed in a tank with 10 g/l of Percy 6340-29 (trade name), with a direct current. After degreasing, a rinse step was performed. Electrolytic graining was carried out in 1% HCl solution, with alternating current of 50 Hz. Rinsing with distilled water followed.

Anodizing was performed in a tank with an electrolyte having a pH of about 2 comprising 400 g/l  $\text{ZnSO}_4 \cdot \text{H}_2\text{O}$  (resulting in about 90-100 g Zn/l), and 30 g/l  $\text{Al}_2\text{SO}_4 \cdot n\text{H}_2\text{O}$ , and 30 g/l  $\text{H}_3\text{BO}_3$ , and with a direct current. The  $\text{H}_3\text{BO}_3$  was added to act as a buffer.

The galvanizing was performed in a separate tank, but with the same electrolytic composition as used for anodising. A rinsing step followed.

From the results of Table 2 it can be seen that all the galvanized sheet material had excellent bonding when tested in a zero bending test. It can also be seen that the line speed, and hence the treatment time, is an important process parameter for the amount of zinc coating on the aluminium sheet. At a line speed of 6 m/min (as in Run 5) the contact times for the various treatment steps were as follows: degreasing 3 s (repeated

four times), graining 5 s, anodizing 5 s, and galvanizing 24 s. The contact times at other line speeds can be calculated from these values.

5            Depending on the local conditions a skilled person  
can therefore find various optima for the process  
parameters depending on the amount of galvanized zinc  
required on the aluminium sheet.

Table 2

Run	Line speed (m/min)	Degrease		Rinse	Graining		Anodizing		Galvanizing		Rinse	Zinc coating (g/m <sup>2</sup> )	Bonding rating
		Temp. (°C)	Current density (A/dm <sup>2</sup> )	Temp. (°C)	Temp. (°C)	Current density (A/dm <sup>2</sup> )	Temp. (°C)	Current density (A/dm <sup>2</sup> )	Temp. (°C)	Current density (A/dm <sup>2</sup> )	Temp. (°C)		
1	4	65	15	51	51	29	52	7.7	50	9.6	52	8.0	1
2	12	65	12.8	46	53	29	52	15.3	51	32	52	5.5	1
3	4	65	12.8	43	54	23	51	15.3	51	32	52	14.5	1
4	2	65	12.8	41	57	23	51	7.7	51	32	52	30.5	1
5	6	66	12.8	37	52	19	52	11.5	52	32	53	26.0	1
6	4	66	12.8	41	48	19	51	11.5	54	32	53	25.5	1
7	2	66	12.8	46	45	19	51	7.7	56	32	52	38.0	1
8	4	66	12.8	49	45	38	52	14.2	58	32	52	41.0	1



CLAIMS

1. A method for applying a layer comprising zinc onto at least one surface of an aluminium or aluminium alloy sheet, comprising the steps of pretreating said surface and applying the layer by electrolytic galvanizing, characterized in that the pretreating step comprises electrochemical graining of said surface.

2. A method according to claim 1 in which the pretreating step comprises anodizing said surface after the electrochemical graining.

3. A method according to claim 1 or 2 in which the electrochemical graining is performed in a solution having a pH less than 3 and in which an alternating current is applied between the sheet and an electrode.

4. A method according to claim 3 in which the electrochemical graining is performed using a current density in the range 5 to 200 A/dm<sup>2</sup>.

5. A method according to claim 3 or 4 in which the alternating current frequency is from 10 to 1000 Hz.

6. A method according to any one of claims 1 to 5 in which the duration of the electrochemical graining is from 0.2 to 60 s.

5 7. A method according to any one of claims 1 to 6 in which the galvanizing is performed using a direct current having a current density of from 10 to 100 A/dm<sup>2</sup>, a bath temperature of from 20 to 70°C, a treatment time of from 5 to 90 s, an electrolyte composition comprising zinc  
10 sulphate having an amount of zinc from 30 to 200 g/l, and an electrolyte pH of less than 4.

8. An aluminium or aluminium alloy sheet having an electrolytically applied layer comprising zinc in which  
15 said layer is adjacent to an aluminium or aluminium alloy surface roughened by electrochemical graining.

9. A method for applying a layer comprising zinc onto at least one surface of an aluminium or aluminium alloy  
20 sheet, comprising the steps of pretreating said surface and applying said layer by electrolytic galvanizing, said pretreating step including applying a preliminary layer comprising zinc onto said surface, characterized in that the application of said preliminary layer comprises (i)  
25 immersing said surface in a zinc-containing alkaline solution, (ii) applying a potential to the sheet to cause

an electrolytic current to flow, and (iii) reversing the polarity of said potential at least once.

5 10. A method according to claim 9 in which the preliminary layer is applied in an amount of from 0.1 to 0.5 g/m<sup>2</sup>.

10 11. A method according to claim 9 or 10 in which said polarity of said potential is reversed at least twice.

12. A method according to claim 11 in which the time interval between each two adjacent polarity reversals is at least 0.5 s.

15 13. A method according to any one of claims 9 to 11 in which during the application of said potential, said sheet is initially an anode and is finally a cathode.

20 14. A method according to any one of claims 9 to 13 in which the current density during the application of said preliminary layer is from 0.5 to 20 A/dm<sup>2</sup>.

25 15. A method according to any one of claims 9 to 14 in which said zinc-containing alkaline solution comprises from 10 to 300 g/l of NaOH and from 2 to 40 g/l of ZnO.

16. A method according to any one of claims 9 to 15 in which said pretreating step includes anodizing said surface after applying said preliminary layer.

5 17. A method according to any one of claims 9 to 16 in which said electrolytic galvanizing is performed with a direct current having a current density of from 10 to 100 A/dm<sup>2</sup>, a bath temperature of from 20 to 70°C, a treatment time of from 5 to 90 s, an electrolyte composition  
10 comprising zinc sulphate in an amount of zinc from 30 to 200 g/l, and an electrolyte pH of less than 4.

18. An aluminium or aluminium alloy sheet having an electrolytically applied layer comprising zinc, produced  
15 by a method in accordance with any one of claims 1 to 7 and 9 to 17.

19. A building sheet comprising an aluminium or aluminium alloy sheet according to claim 8 or 18 in which  
20 the weight of said layer comprising zinc is from 10 to 300 g/m<sup>2</sup>.

20. A building sheet according to claim 19 which is a roofing sheet.

21. A building sheet according to claim 19 or 20 in which the weight of said layer comprising zinc is from 30 to 100 g/m<sup>2</sup>.

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22. An automotive body sheet comprising an aluminium or aluminium alloy sheet according to claim 8 or 18 in which the weight of said layer comprising zinc is from 5 to 40 g/m<sup>2</sup>.

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 97/02329

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 C25D5/44 C25F3/04

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C25D C25F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE WPI Week 7709 Derwent Publications Ltd., London, GB; AN 77-15171y XP002021945 &amp; JP 52 005 630 A (SUMITOMO ELEC. IND. KK) , 17 January 1977 see abstract</p> <p style="text-align: center;">-----</p>	1,8,18

☐ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

### \* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

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"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

21 July 1997

Date of mailing of the international search report

05.08.97

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